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AMENDMENTS TO THE CLAIMS

1. (Amended) A method of treating Polycystic Ovary Syndrome (PCOS) comprising:

identifying a subject suffering from PCOS; and

administering to said subject an effective dose of a composition comprising at least one purified chromium-containing compoundehromium complex.

- 2. (Amended) The method of Claim 1, wherein said <u>purified chromium-containing</u> <u>compoundehromium-complex</u> is selected from the group consisting of chromium picolinate, chromic tripicolinate, chromium nicotinate, chromic polynicotinate, chromium chloride, <u>and</u> chromium histidinate, <u>and chromium yeasts</u>.
- 3. (Original) The method of claim 1, wherein said composition further comprises at least one chelating agent.
- 4. (Original) The method of claim 3, wherein said chelating agent is picolinic acid, nicotinic acid, or both.
- 5. (Amended) The method of claim 4, wherein said <u>purified chromium-containing</u> <u>compoundehromium-complex</u> and said chelating agent are administered in a ratio of between about 1:10 and about 10:1 (w/w).
- 6. (Original) The method of claim 1, further comprising administering a cyclooxygenase inhibitor.
- 7. (Original) The method of claim 6, wherein said cyclooxygenase inhibitor is selected from the group consisting of indomethacin, ibuprofen, acetaminophen, and naproxen.
 - 8. (Original) The method of claim 1, further comprising administering a mucolytic.
 - 9. (Original) The method of claim S. wherein said mucolytic is guaifenesin.
- 10. (Original) The method of claim 1, further comprising administering a salicincontaining herb.
- 11. (Original) The method of claim 10, wherein said salicin-containing herb is selected from the group consisting of *Boswellia serrata* (frankincense), *Betula lenta* (sweet birch), *Betula pubescens* (white birch), *Filipendula ulmaria* (meadowsweet), *Gautheria procumbens* (wintergreens), *Polulus balsamifera*, *Populus jackii* (balm of Gilead) and *Salix alba* (white willow).

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- 12. (Original) The method of claim 1, wherein said effective dose is between about 50 and about 10,000 micrograms.
- 13. (Original) The method of claim 1, wherein said composition is incorporated into a pharmaceutically acceptable carrier selected from the group consisting of a tablet, capsule, microbead, emulsion, powder, granule, suspension, syrup, and elixir.
- 14. (Original) The method of claim 1, wherein said composition is incorporated into a microbead.
- 15. (Amended) The method of claim 14, wherein said microbead is a sugar beadlet or microcrystalline cellulose beadlet and said <u>purified chromium-containing compoundehromium complex</u> is coated on said beadlet.
 - 16. (New) A method of treating Polycystic Ovary Syndrome (PCOS) comprising: identifying a subject suffering from PCOS; and administering to said subject an effective dose of a composition consisting essentially of at least one chromium-containing compound.
- 17. (New) The method of Claim 16, wherein said purified chromium-containing compound is selected from the group consisting of chromium picolinate, chromic tripicolinate, chromium nicotinate, chromic polynicotinate, chromium chloride, and chromium histidinate.
- 18. (New) The method of claim 16, wherein said composition further comprises at least one chelating agent.
- 19. (New) The method of claim 18, wherein said chelating agent is picolinic acid, nicotinic acid, or both.
- 20. (New) The method of claim 19, wherein said purified chromium-containing compound and said chelating agent are administered in a ratio of between about 1:10 and about 10:1 (w/w).
 - 21. (New) A method of treating Polycystic Ovary Syndrome (PCOS) comprising: identifying a subject suffering from PCOS; and administering to said subject an effective dose of a composition comprising at least one synthetic chromium complex.

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- 22. (New) The method of Claim 21, wherein said synthetic chromium complex is selected from the group consisting of chromium picolinate, chromic tripicolinate, chromium nicotinate, chromic polynicotinate, chromium chloride, and chromium histidinate.
- 23. (New) The method of claim 21, wherein said composition further comprises at least one chelating agent.
- 24. (New) The method of claim 23, wherein said chelating agent is picolinic acid, nicotinic acid, or both.
- 25. (New) The method of claim 24, wherein said synthetic chromium complex and said chelating agent are administered in a ratio of between about 1:10 and about 10:1 (w/w).